



Life Science Investor Konference

Copenhagen

October 25 2023

Empowering the body to fight disease

Summary



• Founded 2014, Stockholm-based, listed on Nasdaq First North since 2017

Drug candidate ANXV based on human recombinant protein Annexin A5

• Lead clinical program in RVO with promising signals of effect in Phase 2

Building attractive position in oncology

Active partnering discussions



The team















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Susan Suchdev
Chief Operating Officer

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Chief Financial Officer

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Oncology

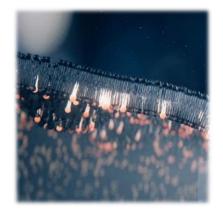
ANXV targets Phosphatidylserine (PS)



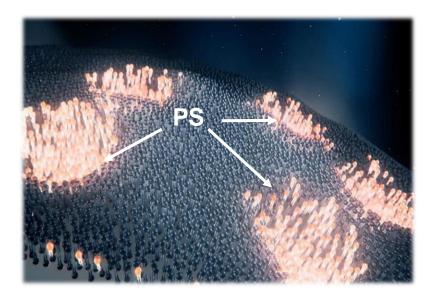
PS externalization is a fingerprint of cell stress/damage/death and triggers multiple pathologic pathways



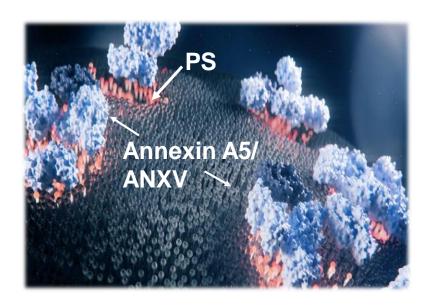
Healthy cell



Healthy cell membrane



Inflammation, RVO, Cancer



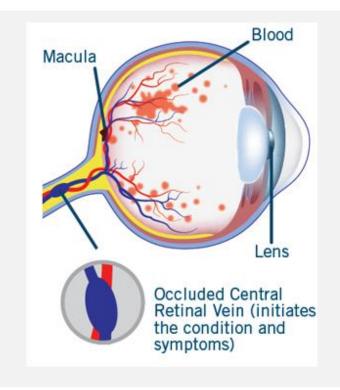
Annexin A5 binds and counteracts PS-signalling

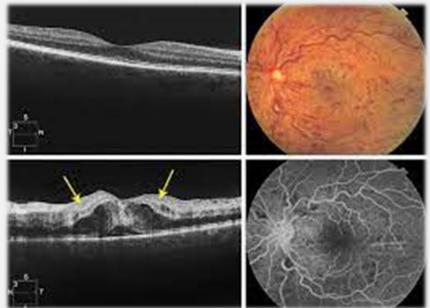


Eye diseases – Retinal Vein Occlusion

Retinal Vein Occlusion – first indication for ANXV^{an}

Pathophysiology and clinical presentation







Aggregation of blood cells in a retinal vein of the eye

Retinal damage swelling, bleeding, cell death, growth of new blood vessels

Impaired vision

Retinal Vein Occlusion – anti-VEGF vs ANXV



Aiming to change current Standard of Care



Retinal Society of Retinal Specialists Brown et al. 2013

- SoC Anti-VEGF treatments are invasive with typically
 6 intravitreal injections
 - e.g. Lucentis® and Eylea®
- Anti-VEGF counteracts complications of RVO retinal swelling and formation of new blood vessels.

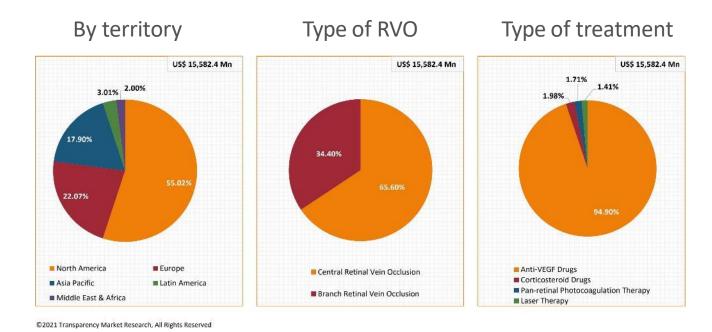


- ANXV short-term intravenous infusion
- ANXV can counteract causative and multiple pathophysiological processes in RVO.

RVO market towards > BUSD 20



Significant and growing market in major territories



Potential ANXV market - conservative scenario

- > 16 M patients globally
- 400.000 new patients per annum in US and EU 5
- App. 150.000 accessible and eligible for ANXV
- Pricing opportunity; current SoC (USD 2000) and consideration of fewer treatments

ANXV in RVO - ongoing Phase 2a



Patient population

Trial Design/Sites

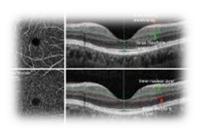
Dosing and Administration

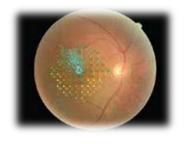
Efficacy, Safety and Adverse Events

Endpoints

- Patients with recently diagnosed RVO, anti VEGF-naïve
- Open label
- Up to 16 RVO patients; 7 sites in the USA
- Dosing schedule 5 days, up to 3 dose levels
- First study with ANXV in a patient population
- Primary-Safety and Tolerability, anti-drug antibodies
- Secondary pharmacokinetic data
- Exploratory anatomical and functional investigations, PS on erythrocytes











BCVA

UWF-FA

OCT-A

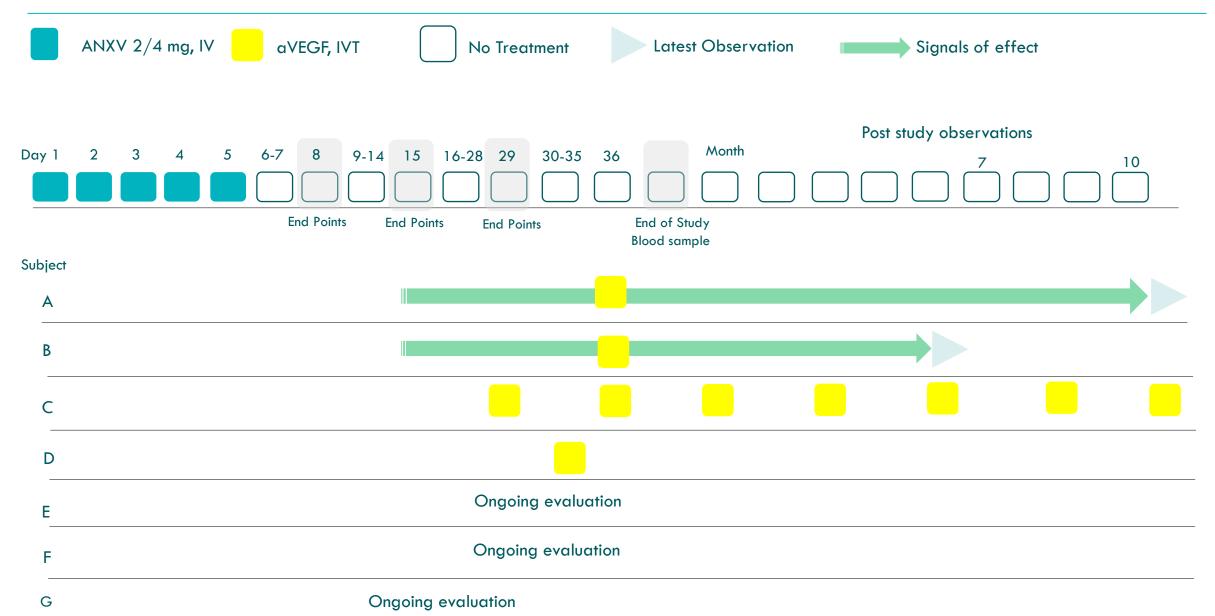
MMP

rescue with anti-VEGF

PS target
ClinTrials NCT05532735

ANN 004 - Open Label Study Design



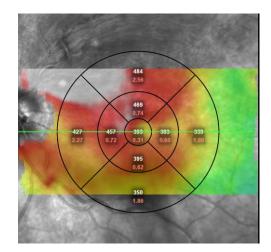


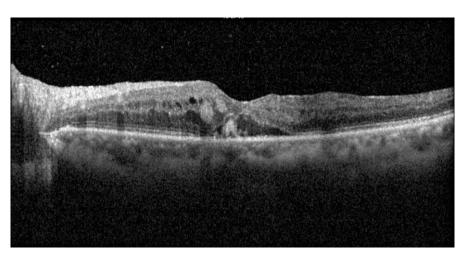
No swelling and stable visual acuity at month 10

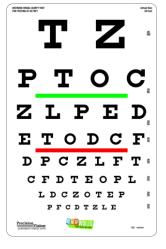


Subject A, "ANXV + one dose of aVEGF, a happy, highly unusual and potentially cured patient" PI

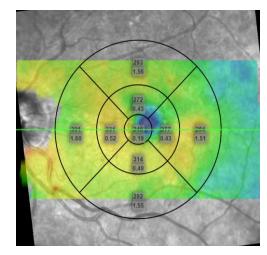
Day 0

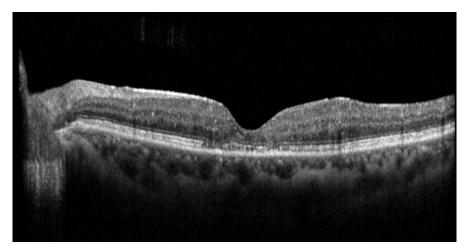


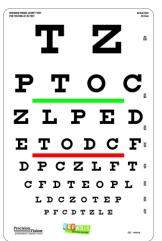




Month 6











ANXV in RVO - market plan for partnering discussions

Potential for Phase 2b and effective Phase 3 trials

- Phase 2b
 - Opportunity to use reduced need for anti-VEGF injections
 - Preliminary primary endpoint for significance:
 - reduced no of anti-VEGF injections at 6 months
 - Phase 3 guiding secondary endpoints

- Phase 3
 - Regulatory advice (FDA) regarding approvable endpoints for marketing authorisation
 - BCVA is standard based on anti-VEGF approvals
 - BCVA in combination with reduced anti-VEGF requirements?

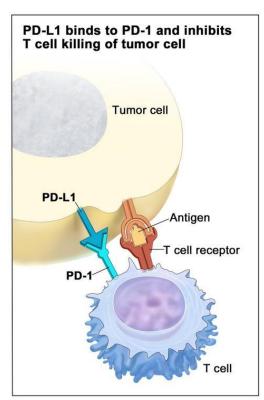


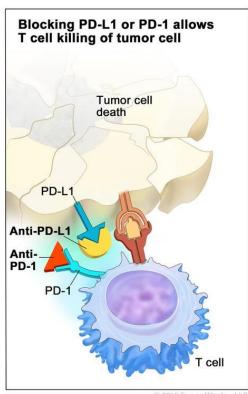
Oncology

Modern principles to treat cancers



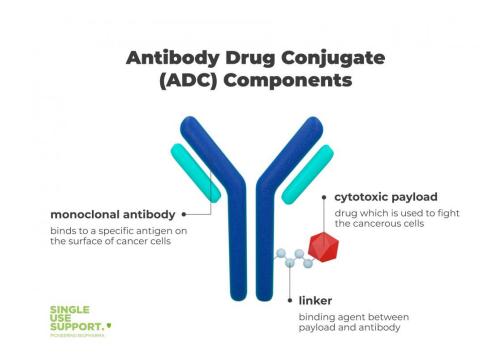
Immune checkpoint inhibition





© 2015 Terese Winslow LLC

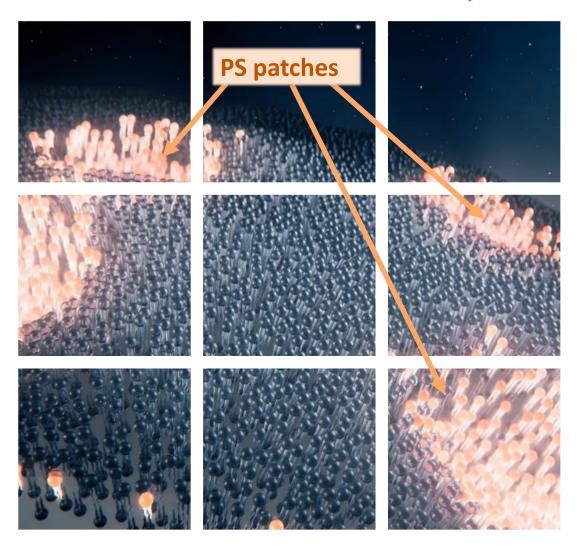
Targeted delivery of chemotherapy



Tumors externalize PS to hide from immune system

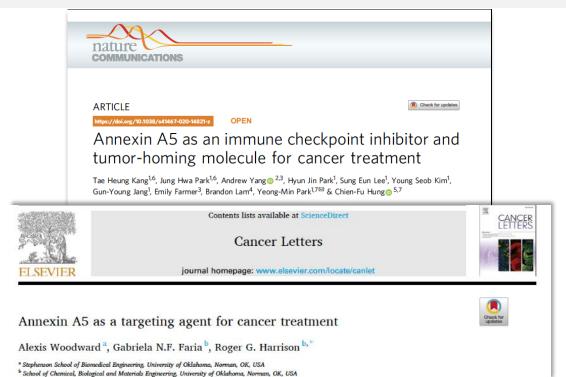


PS is a fundamental immune checkpoint



Phosphatidylserine (PS) is

- a universal marker for cancer
- used by cancer cells to survive by inactivating immune cells
- proposed as an "Achilles heal of cancers" potential for therapeutic purposes



Immune checkpoint inhibition market



Intense development, marketing and sales of immunoncology drugs for cancer patients

Keytruda (PD-1, Merck)

• App. **BUSD 21** annual sales

Opdivo (PD-1, BMS)

App. BUSD 8 annual sales

Yervoy (CTL-4, BMS)

App. MUSD 500 annual sales

Immuno-oncology
Checkpoint inhibition

ANXV market opportunity in immuno-oncology across 3 indication examples if ANXV becomes 2nd line IO

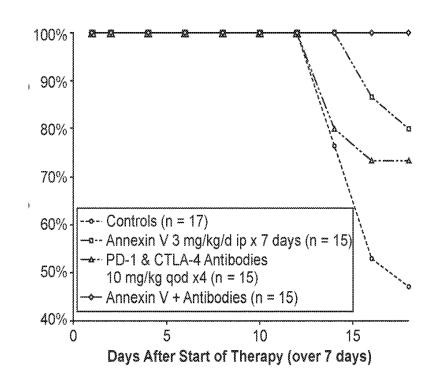
Cancer indication	IO Market size 2026 forecast	Failure rate 1 st line IO %	Eligibility 2 nd line IO/ANXV %	ANXV market value Estimate BUSD
	BUSD			
NSCLC	43	60	40	10.3
Melanoma	4.2	30	32	0.6
Head & neck	2.9	30	30	0.61

Improvement of survival in animal models



- Robust animal data from independent labs
- Anti-tumor effect as good as than current marketed CPIs

Survival, % of animals



ANXV

Immuno-oncology
Checkpoint inhibition

Lethal tumor challenge, lung cancer

- 7 days of Annexin A5 treatment
- ANXV as mono- or combination-therapy allowed up to 100% survival

Blankenberg, US Patent

Annexin A5 – a carrier of validated chemotherapy drugs



ANXV has potentially competitive advantages

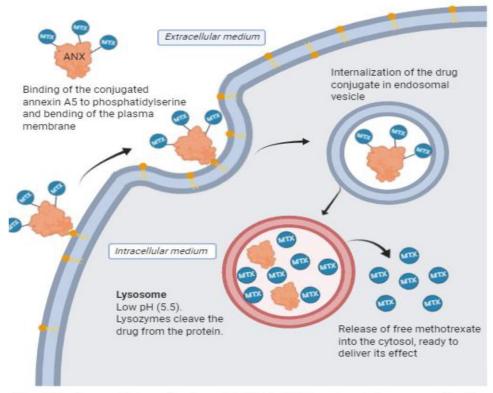


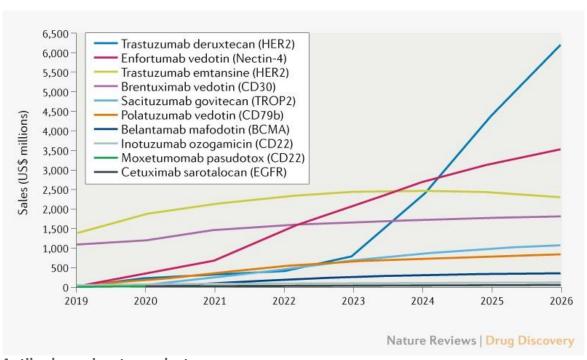
Fig. 6 Binding and internalization of ANXA5-MTX conjugate in cancer cells. The drug conjugate binds to externalized phosphatidylserine (yellow) of cancer cells, internalizes and releases active methotrexate in the cytosol. The figure was drawn using the software © *BioRender 2020*.

Targeted delivery of chemotherapy

ANXV drug conjugate can enter a substantial market



Currently marketed products target specific receptors for specific cancers ANXV-drug conjugate may have broader applications



Market reports confirm value proposition					
ADC Market Size in 2021	USD 4.2 Billion				
Projected Market Size in 2028	USD 13.2 Billion				
CAGR Growth Rate 2022- 2028	18.9%				

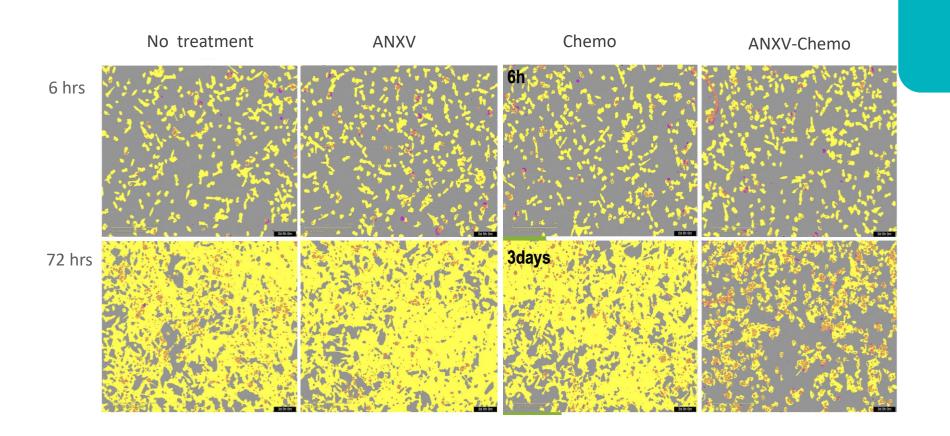
Antibody-conjugate products

- Kadcycla, (Roche, yellow) Her2 positive breast cancers
- Enhertu (Daiichi-Sankyo/AstraZeneca, blue), Her2 positive breast cancers
- Padcev (Seagen/Astellas, rose) Nectin 4 positive urothelial cancers
- Adectris (Seagen/Takeda, purple), CD30 positive Hodgkin lymphomas

ANXV-chemo kills PSpos aggressive cancer cells



In vitro Proof of Principle – triple negative breastcancer cells



ANXV-chemo
Targeted delivery
of chemotherapy

Summary



Drug candidate ANXV based on human recombinant protein Annexin A5

Broad therapeutic potential

Derisked program in clinical stage

Lead clinical program in RVO with promising signals of effect in Phase 2

Building attractive position in oncology

Rights issue of 35.6 MSEK, 100% guaranteed

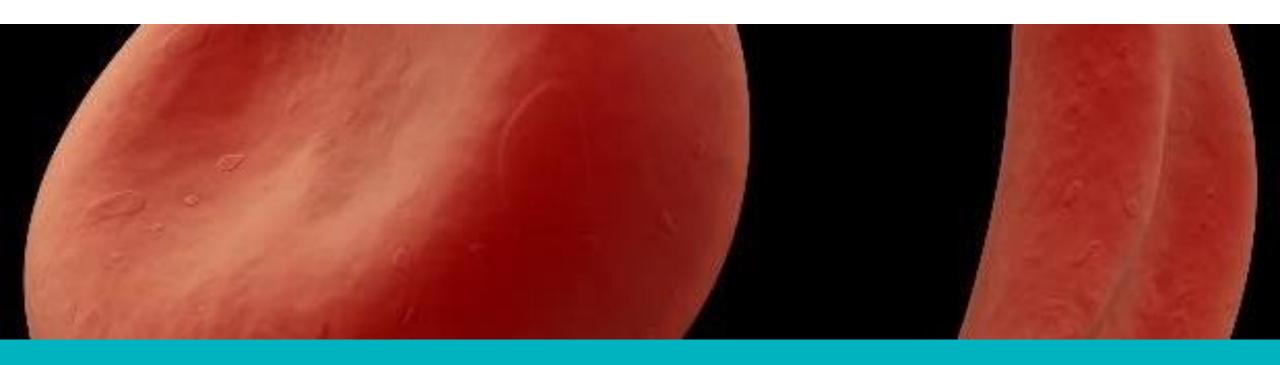


- The Structure
 - 35.6 MSEK gross
 - 100% guaranteed by main shareholders
 - Subscription price 0.20 SEK
 - 11:10 For each 10 existing shares, shareholders can subscribe for 11 new shares
- The Time plan:

24 November 2023	Extraordinary General Meeting	
24 November 2023	Last day of trading in the share including the	
	right to receive subscription rights	
27 November 2023	First day of trading in the share excluding the	
	right to receive subscription rights	
28 November 2023	Record date for participating in the Rights Issue	
28 November 2023	Estimated date for the publication of the	
	Prospectus	
30 November – 11 December 2023	Trading in subscription rights	
30 November – 14 December 2023	Subscription period	
30 November 2023 – Until the Rights	Trading in paid subscribed shares (Sw. "BTA")	
Issue is registered with the Swedish		
Companies Registration Office		
Around 18 December 2023	Estimated publication of the outcome in the	
	Rights Issue	

https://www.annexinpharma.com





Leveraging therapeutic opportunities with ANXV